

# **Inhibitory effects of an aqueous extract of *Clitoria ternatea* flower on $\alpha$ -glucosidase during in-vitro wheat starch digestion**

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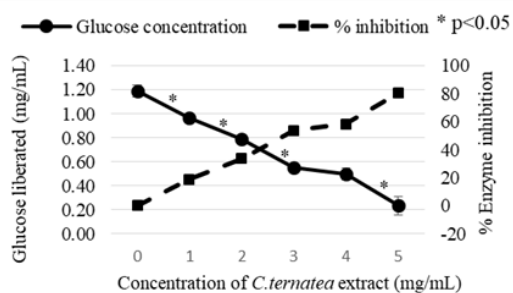
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**Inhibitory effects of an aqueous extract of *Clitoria ternatea* flower on  $\alpha$ -glucosidase during *in-vitro* wheat starch digestion.** By C. Ronald, B.S. Chu, *Division of Food and Drink, School of Applied Sciences, Abertay University, Bell Street, Dundee, DD1 1HG*

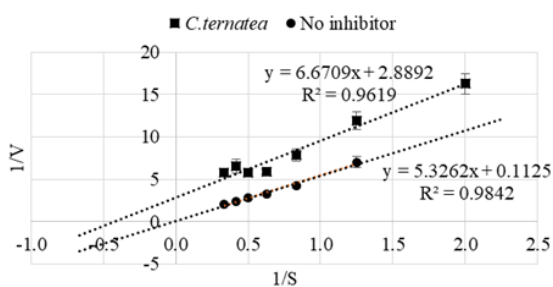
Phenolic compounds in plants inhibit  $\alpha$ -glucosidase, reducing carbohydrate absorption and contributing to anti-diabetic activity<sup>(1)</sup>, particularly anthocyanins from coloured fruit and vegetables such as berries<sup>(2)</sup> and black and purple carrots<sup>(3)</sup>. The aim of this study was to evaluate inhibition behaviours of an extract from the blue *C. ternatea* flower against  $\alpha$ -glucosidase during *in-vitro* wheat starch digestion i.e. the potential of the flower extract to modulate starch digestion in a bid to reduce postprandial hyperglycaemic response.

Inhibition of an aqueous extract of dried *C. ternatea* flowers, at different concentrations, against  $\alpha$ -glucosidase was determined according to the method of Sui *et al.*<sup>(4)</sup> with minor modifications. In Eppendorf tubes, at each extract concentration, aliquots were mixed with the extract, 0.5 mg/mL  $\alpha$ -glucosidase, 40 mg/L calcium chloride, and distilled water to 712  $\mu$ L, then incubated at 37°C for 15 min for the enzyme to interact with the extract. Aliquots were made up to a 750  $\mu$ L reaction solution with 1 mg/mL gelatinised starch solution and incubated at 37°C for 5 min to simulate starch digestion. Exactly 100  $\mu$ L 3,5-dinitrosalicylic acid reagent solution was added to each tube, heated for 10 min in boiling water, then cooled for 10 min in an ice bath; glucose concentration was then measured. Experiments were repeated with *C. ternatea* extract at 3 mg/mL and differing starch concentrations; 0.5-2 mg/mL. All experiments were repeated in triplicate.

Results showed decreases in glucose liberated and increases in %inhibition, after 5 min digestion, as concentrations of *C. ternatea* extract increased. Around 80% inhibition was reached at 5 mg/mL *C. ternatea* extract. Significant statistical differences ( $p < 0.001$ ) were found between the means of *C. ternatea* extract groups as determined by one-way ANOVA. Tukey post hoc tests showed that mean glucose liberated between each concentration was statistically significantly lower as concentration increased ( $p < 0.05$ ), except between concentrations 3 and 4 mg/mL, where the difference found was not statistically significant ( $p = 0.421$ ). Inhibition activity of *C. ternatea* extract against  $\alpha$ -glucosidase was found to be uncompetitive with  $K_M = 2.309$  mg/mL,  $V_{MAX} = 0.346$  mgmin<sup>-1</sup> and  $IC_{50} = 2.315$  mg/mL.



**Fig. 1.** Effect of *C. ternatea* extract on % inhibition of  $\alpha$ -glucosidase.



**Fig. 2.** Lineweaver-Burk plot; with and without *C. ternatea* extract to assess type of inhibition against  $\alpha$ -glucosidase.

These results suggest that *C. ternatea* extract has potential to be utilised to prepare functional foods and/or nutraceuticals to help control postprandial hyperglycaemic response therefore, could be used to treat, and reduce the risk of developing, Type 2 diabetes.

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